

**What is claimed is:**

1. An antagonist that inhibits or an agonist that activates an activity a polypeptide selected from the group consisting of: a polypeptide comprising an amino acid sequence which is at least 90% identical to the amino acid sequence of SEQ ID NO:2, and a polypeptide comprising an amino acid sequence as set forth in SEQ ID NO:2, wherein said activity is selected from the group consisting of: NADPH-dependent reduction of acetoacetyl-acyl carrier protein (ACP) to generate  $\beta$ -hydroxyacyl-ACP; deprotonation of a group leading to a diminution in  $k_{cat}$ ; deprotonation of a general acid responsible for donating a proton to the carbonyl oxygen during its reduction; binding of an ionizable group putatively binding the pyrophosphate bridge of NADPH; catalysis involving a lysine residue as a general acid; a conformational change inducing formation a low-barrier hydrogen bond (LBHB) in ketone reduction mechanism; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB between Tyr157 and Lys161; energy provided from Tyr157 and Lys161; energy from forming an LBHB between Tyr157 and Lys161 facilitating proton transfer from Lys157 to the carbonyl oxygen; proton transfer from Lys157 to carbonyl oxygen; formation of an LBHB between Tyr157 and an Asp residue; strengthening of the role of Tyr157 in facilitating general acid catalysis; strengthening of the role of Tyr157 in facilitating general acid catalysis by Lys161; compression of active site; compression of active site resulting in the formation of a LBHB facilitating proton transfer to the carbonyl oxygen; hydride transfer from NADPH proceeding proton transfer from the Lys residue; formation of an anionic, tetrahedral reaction intermediate; formation of a charge-stabilized intermediate by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2); and hydride transfer from NADPH proceeding proton transfer from the Lys residue, with an anionic, tetrahedral reaction intermediate (1) being formed, that is potentially charge-stabilized by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2).

2. A method for the treatment of an individual having need to inhibit or activate Fab G polypeptide comprising the steps of: administering to the individual an antibacterially effective amount of an antagonist that inhibits or an agonist that activates an activity of a polypeptide selected from the group consisting of: a polypeptide comprising an amino acid sequence which is at least 90% identical to the amino acid sequence of SEQ ID NO:2, and a polypeptide comprising an amino acid sequence as set forth in SEQ ID NO:2, wherein said

activity is selected from the group consisting of: NADPH-dependent reduction of acetoacetyl-acyl carrier protein (ACP) to generate  $\beta$ -hydroxyacyl-ACP; deprotonation of a group leading to a diminution in  $k_{cat}$ ; deprotonation of a general acid responsible for donating a proton to the carbonyl oxygen during its reduction; binding of an ionizable group putatively binding the pyrophosphate bridge of NADPH; catalysis involving a lysine residue as a general acid; a conformational change inducing formation a low-barrier hydrogen bond (LBHB) in ketone reduction mechanism; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB between Tyr157 and Lys161; energy provided from Tyr157 and Lys161; energy from forming an LBHB between Tyr157 and Lys161 facilitating proton transfer from Lys157 to the carbonyl oxygen; proton transfer from Lys157 to carbonyl oxygen; formation of an LBHB between Tyr157 and an Asp residue; strengthening of the role of Tyr157 in facilitating general acid catalysis; strengthening of the role of Tyr157 in facilitating general acid catalysis by Lys161; compression of active site; compression of active site resulting in the formation of a LBHB facilitating proton transfer to the carbonyl oxygen; hydride transfer from NADPH proceeding proton transfer from the Lys residue; formation of an anionic, tetrahedral reaction intermediate; formation of a charge-stabilized intermediate by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2); and hydride transfer from NADPH proceeding proton transfer from the Lys residue, with an anionic, tetrahedral reaction intermediate (1) being formed, that is potentially charge-stabilized by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2).

3. A method for the treatment of an individual infected with a bacteria comprising the steps of administering to the individual an antibacterially effective amount of an antagonist that inhibits or an agonist that activates an activity of a polypeptide selected from the group consisting of: a polypeptide comprising an amino acid sequence which is at least 90% identical to the amino acid sequence of SEQ ID NO:2, and a polypeptide comprising an amino acid sequence as set forth in SEQ ID NO:2, wherein said activity is selected from the group consisting of: NADPH-dependent reduction of acetoacetyl-acyl carrier protein (ACP) to generate  $\beta$ -hydroxyacyl-ACP; deprotonation of a group leading to a diminution in  $k_{cat}$ ; deprotonation of a general acid responsible for donating a proton to the carbonyl oxygen during its reduction; binding of an ionizable group putatively binding the pyrophosphate bridge of NADPH; catalysis involving a lysine residue as a general acid; a conformational change

inducing formation a low-barrier hydrogen bond (LBHB) in ketone reduction mechanism; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB between Tyr157 and Lys161; energy provided from Tyr157 and Lys161; energy from forming an LBHB between Tyr157 and Lys161 facilitating proton transfer from Lys157 to the carbonyl oxygen; proton transfer from Lys157 to carbonyl oxygen; formation of an LBHB between Tyr157 and an Asp residue; strengthening of the role of Tyr157 in facilitating general acid catalysis; strengthening of the role of Tyr157 in facilitating general acid catalysis by Lys161; compression of active site; compression of active site resulting in the formation of a LBHB facilitating proton transfer to the carbonyl oxygen; hydride transfer from NADPH proceeding proton transfer from the Lys residue; formation of an anionic, tetrahedral reaction intermediate; formation of a charge-stabilized intermediate by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2); and hydride transfer from NADPH proceeding proton transfer from the Lys residue, with an anionic, tetrahedral reaction intermediate (1) being formed, that is potentially charge-stabilized by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2).

4. The method of claim 3 wherein said bacteria is selected from the group consisting of a member of the genus *Staphylococcus*, *Staphylococcus aureus*, a member of the genus *Streptococcus*, and *Streptococcus pneumoniae*.

5. A method for the treatment of an individual having need to inhibit or activate Fab G polypeptide comprising the steps of administering to the individual an antibacterially effective amount of an antagonist that inhibits or an agonist that activates an activity of Fab G selected from the group consisting of: NADPH-dependent reduction of acetoacetyl-acyl carrier protein (ACP) to generate  $\beta$ -hydroxyacyl-ACP; deprotonation of a group leading to a diminution in  $k_{\text{cat}}$ ; deprotonation of a general acid responsible for donating a proton to the carbonyl oxygen during its reduction; binding of an ionizable group putatively binding the pyrophosphate bridge of NADPH; catalysis involving a lysine residue as a general acid; a conformational change inducing formation a low-barrier hydrogen bond (LBHB) in ketone reduction mechanism; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB between Tyr157 and Lys161; energy provided from Tyr157 and Lys161; energy from forming an LBHB between Tyr157 and Lys161 facilitating proton transfer from

Lys157 to the carbonyl oxygen; proton transfer from Lys157 to carbonyl oxygen; formation of an LBHB between Tyr157 and an Asp residue; strengthening of the role of Tyr157 in facilitating general acid catalysis; strengthening of the role of Tyr157 in facilitating general acid catalysis by Lys161; compression of active site; compression of active site resulting in the formation of a LBHB facilitating proton transfer to the carbonyl oxygen; hydride transfer from NADPH proceeding proton transfer from the Lys residue; formation of an anionic, tetrahedral reaction intermediate; formation of a charge-stabilized intermediate by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2); and hydride transfer from NADPH proceeding proton transfer from the Lys residue (Figure 5), with an anionic, tetrahedral reaction intermediate (1) being formed, that is potentially charge-stabilized by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2).

6. A method for the treatment of an individual infected with a bacteria comprising the steps of administering to the individual an antibacterially effective amount of an antagonist that inhibits or an agonist that activates that activates an activity of Fab G selected from the group consisting of: NADPH-dependent reduction of acetoacetyl-acyl carrier protein (ACP) to generate  $\beta$ -hydroxyacyl-ACP; deprotonation of a group leading to a diminution in  $k_{cat}$ ; deprotonation of a general acid responsible for donating a proton to the carbonyl oxygen during its reduction; binding of an ionizable group putatively binding the pyrophosphate bridge of NADPH; catalysis involving a lysine residue as a general acid; a conformational change inducing formation a low-barrier hydrogen bond (LBHB) in ketone reduction mechanism; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB between Tyr157 and Lys161; energy provided from Tyr157 and Lys161; energy from forming an LBHB between Tyr157 and Lys161 facilitating proton transfer from Lys157 to the carbonyl oxygen; proton transfer from Lys157 to carbonyl oxygen; formation of an LBHB between Tyr157 and an Asp residue; strengthening of the role of Tyr157 in facilitating general acid catalysis; strengthening of the role of Tyr157 in facilitating general acid catalysis by Lys161; compression of active site; compression of active site resulting in the formation of a LBHB facilitating proton transfer to the carbonyl oxygen; hydride transfer from NADPH proceeding proton transfer from the Lys residue; formation of an anionic, tetrahedral reaction intermediate; formation of a charge-stabilized intermediate by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2); and hydride transfer from NADPH proceeding proton

transfer from the Lys residue, with an anionic, tetrahedral reaction intermediate (1) being formed, that is potentially charge-stabilized by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2).

7. The method of claim 6 wherein said bacteria is selected from the group consisting of: a member of the genus *Staphylococcus*, *Staphylococcus aureus*, a member of the genus *Streptococcus*, and *Streptococcus pneumoniae*.

8. A method for the treatment of an individual infected by *Streptococcus pneumoniae* comprising the steps of administering to the individual an antibacterially effective amount of an antagonist that inhibits or antagonist that activates an activity of *Streptococcus pneumoniae* Fab G selected from the group consisting of: NADPH-dependent reduction of acetoacetyl-acyl carrier protein (ACP) to generate  $\beta$ -hydroxyacyl-ACP; deprotonation of a group leading to a diminution in  $k_{cat}$ ; deprotonation of a general acid responsible for donating a proton to the carbonyl oxygen during its reduction; binding of an ionizable group putatively binding the pyrophosphate bridge of NADPH; catalysis involving a lysine residue as a general acid; a conformational change inducing formation a low-barrier hydrogen bond (LBHB) in ketone reduction mechanism; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB between Tyr157 and Lys161; energy provided from Tyr157 and Lys161; energy from forming an LBHB between Tyr157 and Lys161 facilitating proton transfer from Lys157 to the carbonyl oxygen; proton transfer from Lys157 to carbonyl oxygen; formation of an LBHB between Tyr157 and an Asp residue; strengthening of the role of Tyr157 in facilitating general acid catalysis; strengthening of the role of Tyr157 in facilitating general acid catalysis by Lys161; compression of active site; compression of active site resulting in the formation of a LBHB facilitating proton transfer to the carbonyl oxygen; hydride transfer from NADPH proceeding proton transfer from the Lys residue; formation of an anionic, tetrahedral reaction intermediate; formation of a charge-stabilized intermediate by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2); and hydride transfer from NADPH proceeding proton transfer from the Lys residue, with an anionic, tetrahedral reaction intermediate (1) being formed, that is potentially charge-stabilized by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2).

9. An antagonist that inhibits an activity of a polypeptide selected from the group consisting of: a polypeptide comprising an amino acid sequence which is at least 90% identical

to the amino acid sequence of SEQ ID NO:1, and a polypeptide comprising an amino acid sequence as set forth in SEQ ID NO:1, wherein said activity is selected from the group consisting of: NADPH-dependent reduction of acetoacetyl-acyl carrier protein (ACP) to generate  $\beta$ -hydroxyacyl-ACP; deprotonation of a group leading to a diminution in  $k_{cat}$ ; deprotonation of a general acid responsible for donating a proton to the carbonyl oxygen during its reduction; binding of an ionizable group putatively binding the pyrophosphate bridge of NADPH; catalysis involving a lysine residue as a general acid; a conformational change inducing formation a low-barrier hydrogen bond (LBHB) in ketone reduction mechanism; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB between Tyr157 and Lys161; energy provided from Tyr157 and Lys161; energy from forming an LBHB between Tyr157 and Lys161 facilitating proton transfer from Lys157 to the carbonyl oxygen; proton transfer from Lys157 to carbonyl oxygen; formation of an LBHB between Tyr157 and an Asp residue; strengthening of the role of Tyr157 in facilitating general acid catalysis; strengthening of the role of Tyr157 in facilitating general acid catalysis by Lys161; compression of active site; compression of active site resulting in the formation of a LBHB facilitating proton transfer to the carbonyl oxygen; hydride transfer from NADPH proceeding proton transfer from the Lys residue; formation of an anionic, tetrahedral reaction intermediate; formation of a charge-stabilized intermediate by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2); and hydride transfer from NADPH proceeding proton transfer from the Lys residue, with an anionic, tetrahedral reaction intermediate (1) being formed, that is potentially charge-stabilized by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2).

10. A method for the treatment of an individual having need to inhibit Fab G polypeptide comprising the steps of administering to the individual an antibacterially effective amount of an antagonist that inhibits an activity of a polypeptide selected from the group consisting of: a polypeptide comprising an amino acid sequence which is at least 90% identical to the amino acid sequence of SEQ ID NO:1, and a polypeptide comprising an amino acid sequence as set forth in SEQ ID NO:1, wherein said activity is selected from the group consisting of: NADPH-dependent reduction of acetoacetyl-acyl carrier protein (ACP) to generate  $\beta$ -hydroxyacyl-ACP; deprotonation of a group leading to a diminution in  $k_{cat}$ ; deprotonation of a general acid responsible for donating a proton to the carbonyl oxygen during

its reduction; binding of an ionizable group putatively binding the pyrophosphate bridge of NADPH; catalysis involving a lysine residue as a general acid; a conformational change inducing formation a low-barrier hydrogen bond (LBHB) in ketone reduction mechanism; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB between Tyr157 and Lys161; energy provided from Tyr157 and Lys161; energy from forming an LBHB between Tyr157 and Lys161 facilitating proton transfer from Lys157 to the carbonyl oxygen; proton transfer from Lys157 to carbonyl oxygen; formation of an LBHB between Tyr157 and an Asp residue; strengthening of the role of Tyr157 in facilitating general acid catalysis; strengthening of the role of Tyr157 in facilitating general acid catalysis by Lys161; compression of active site; compression of active site resulting in the formation of a LBHB facilitating proton transfer to the carbonyl oxygen;

hydride transfer from NADPH preceding proton transfer from the Lys residue; formation of an anionic, tetrahedral reaction intermediate; formation of a charge-stabilized intermediate by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2); and hydride transfer from NADPH preceding proton transfer from the Lys residue, with an anionic, tetrahedral reaction intermediate (1) being formed, that is potentially charge-stabilized by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2).

11. A method for inhibiting an activity of Fab G polypeptide comprising the steps of contacting a composition comprising said polypeptide with an effective amount of an antagonist that inhibits an activity of Fab G, wherein said activity is selected from the group consisting of: NADPH-dependent reduction of acetoacetyl-acyl carrier protein (ACP) to generate  $\beta$ -hydroxyacyl-ACP; deprotonation of a group leading to a diminution in  $k_{cat}$ ; deprotonation of a general acid responsible for donating a proton to the carbonyl oxygen during its reduction; binding of an ionizable group putatively binding the pyrophosphate bridge of NADPH; catalysis involving a lysine residue as a general acid; a conformational change inducing formation a low-barrier hydrogen bond (LBHB) in ketone reduction mechanism; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB between Tyr157 and Lys161; energy provided from Tyr157 and Lys161; energy from forming an LBHB between Tyr157 and Lys161 facilitating proton transfer from Lys157 to the carbonyl oxygen; proton transfer from Lys157 to carbonyl oxygen; formation of an LBHB between

Tyr157 and an Asp residue; strengthening of the role of Tyr157 in facilitating general acid catalysis; strengthening of the role of Tyr157 in facilitating general acid catalysis by Lys161; compression of active site; compression of active site resulting in the formation of a LBHB facilitating proton transfer to the carbonyl oxygen; hydride transfer from NADPH proceeding proton transfer from the Lys residue; formation of an anionic, tetrahedral reaction intermediate; formation of a charge-stabilized intermediate by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2); and hydride transfer from NADPH proceeding proton transfer from the Lys residue, with an anionic, tetrahedral reaction intermediate (1) being formed, that is potentially charge-stabilized by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2).

12. A method for inhibiting an activity of Fab G, wherein said activity is selected from the group consisting of: NADPH-dependent reduction of acetoacetyl-acyl carrier protein (ACP) to generate  $\beta$ -hydroxyacyl-ACP; deprotonation of a group leading to a diminution in  $k_{cat}$  (Figure 1A); deprotonation of a general acid responsible for donating a proton to the carbonyl oxygen during its reduction; binding of an ionizable group putatively binding the pyrophosphate bridge of NADPH; catalysis involving a lysine residue as a general acid; a conformational change inducing formation a low-barrier hydrogen bond (LBHB) in ketone reduction mechanism; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB between Tyr157 and Lys161; energy provided from Tyr157 and Lys161; energy from forming an LBHB between Tyr157 and Lys161 facilitating proton transfer from Lys157 to the carbonyl oxygen; proton transfer from Lys157 to carbonyl oxygen; formation of an LBHB between Tyr157 and an Asp residue; strengthening of the role of Tyr157 in facilitating general acid catalysis; strengthening of the role of Tyr157 in facilitating general acid catalysis by Lys161; compression of active site; compression of active site resulting in the formation of a LBHB facilitating proton transfer to the carbonyl oxygen; hydride transfer from NADPH proceeding proton transfer from the Lys residue; formation of an anionic, tetrahedral reaction intermediate; formation of a charge-stabilized intermediate by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2); and hydride transfer from NADPH proceeding proton transfer from the Lys residue, with an anionic, tetrahedral reaction intermediate (1) being formed, that is potentially charge-stabilized by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2).

13. The method of claim 12 wherein said bacteria is selected from the group consisting of: a member of the genus *Staphylococcus*, *Staphylococcus aureus*, a member of the genus *Streptococcus*, and *Streptococcus pneumoniae*.

14. A method for inhibiting a growth of bacteria comprising the steps of contacting a composition comprising bacteria with an antibacterially effective amount of an antagonist that inhibits an activity of Fab G, wherein said activity is selected from the group consisting of: NADPH-dependent reduction of acetoacetyl-acyl carrier protein (ACP) to generate  $\beta$ -hydroxyacyl-ACP; deprotonation of a group leading to a diminution in  $k_{cat}$ ; deprotonation of a general acid responsible for donating a proton to the carbonyl oxygen during its reduction; binding of an ionizable group putatively binding the pyrophosphate bridge of NADPH; catalysis involving a lysine residue as a general acid; a conformational change inducing formation a low-barrier hydrogen bond (LBHB) in ketone reduction mechanism; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB between Tyr157 and Lys161; energy provided from Tyr157 and Lys161; energy from forming an LBHB between Tyr157 and Lys161 facilitating proton transfer from Lys157 to the carbonyl oxygen; proton transfer from Lys157 to carbonyl oxygen; formation of an LBHB between Tyr157 and an Asp residue; strengthening of the role of Tyr157 in facilitating general acid catalysis; strengthening of the role of Tyr157 in facilitating general acid catalysis by Lys161; compression of active site; compression of active site resulting in the formation of a LBHB facilitating proton transfer to the carbonyl oxygen; hydride transfer from NADPH proceeding proton transfer from the Lys residue; formation of an anionic, tetrahedral reaction intermediate; formation of a charge-stabilized intermediate by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2); and hydride transfer from NADPH proceeding proton transfer from the Lys residue, with an anionic, tetrahedral reaction intermediate (1) being formed, that is potentially charge-stabilized by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2).

15. The method of claim 14 wherein said bacteria is selected from the group consisting of: a member of the genus *Staphylococcus*, *Staphylococcus aureus*, a member of the genus *Streptococcus*, and *Streptococcus pneumoniae*.

16. A method for inhibiting a Fab G polypeptide comprising the steps of contacting a composition comprising bacteria with an antibacterially effective amount of an antagonist that inhibits an activity of Fab G, wherein said activity is selected from the group consisting of:

NADPH-dependent reduction of acetoacetyl-acyl carrier protein (ACP) to generate  $\beta$ -hydroxyacyl-ACP; deprotonation of a group leading to a diminution in  $k_{cat}$ ; deprotonation of a general acid responsible for donating a proton to the carbonyl oxygen during its reduction; binding of an ionizable group putatively binding the pyrophosphate bridge of NADPH; catalysis involving a lysine residue as a general acid; a conformational change inducing formation a low-barrier hydrogen bond (LBHB) in ketone reduction mechanism; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB; a conformational change upon acetoacetyl-CoA binding resulting in formation of an LBHB between Tyr157 and Lys161; energy provided from Tyr157 and Lys161; energy from forming an LBHB between Tyr157 and Lys161 facilitating proton transfer from Lys157 to the carbonyl oxygen; proton transfer from Lys157 to carbonyl oxygen; formation of an LBHB between Tyr157 and an Asp residue; strengthening of the role of Tyr157 in facilitating general acid catalysis; strengthening of the role of Tyr157 in facilitating general acid catalysis by Lys161; compression of active site; compression of active site resulting in the formation of a LBHB facilitating proton transfer to the carbonyl oxygen; hydride transfer from NADPH proceeding proton transfer from the Lys residue; formation of an anionic, tetrahedral reaction intermediate; formation of a charge-stabilized intermediate by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2); and hydride transfer from NADPH proceeding proton transfer from the Lys residue, with an anionic, tetrahedral reaction intermediate (1) being formed, that is potentially charge-stabilized by protonated Lys group prior to proton transfer to form the  $\beta$ -hydroxy-keto product (2).

17. The method of claim 16 wherein said bacteria is selected from the group consisting of: a member of the genus *Staphylococcus*, *Staphylococcus aureus*, a member of the genus *Streptococcus*, and *Streptococcus pneumoniae*.